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| 10/648,536 | 08/25/2003 | Robert Owen Lockerbie | B0175-US02 | 4649 |
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| CaridianBCT, Inc. Mail Stop: 810 1F2 10811 WEST COLLINS AVE LAKEWOOD, CO 80215 | | | EXAMINER LEE, JAE W | |
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Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

DETAILED ACTION

Application status

In response to the previous Office action, a non-final rejection (mailed on 05/20/2009), Applicants filed an amendment to claims on 08/06/2009.

Maintained Rejections

Claim Rejections - 35 U.S.C. § 103

The following is a quotation of 35 U.S.C. § 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

Claims 12-15, 17-19 and 22 are rejected under 35 U.S.C. § 103(a) as being unpatentable over Goodrich et al.¹ (USPN 6,258,577) in view of Joshi PC (Comparison of the DNA-damaging property of photosensitised riboflavin via singlet oxygen (1O₂) and superoxide radical O₂⁻. Mechanisms, Toxicol Lett. 1985, 26(2-3):211-7).

Applicants argue that all of the elements of Applicants claims were not known from either the Goodrich or Joshi references. There is no disclosure in Goodrich of using riboflavin and light (at any wavelength) to inactivate white blood cells. There is also no disclosure of using riboflavin and light to cause damage to the nucleic acids of white blood cells and substantially maintaining the damage to the nucleic acids of white

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blood cells. Thirdly, and as the Examiner admits, Goodrich does not disclose the specific use of UVB light to inactivate white blood cells. The Joshi reference does not cure the deficiencies of the Goodrich reference. The Joshi reference teaches that riboflavin generates singlet oxygen and superoxide anion radicals upon exposure to UVB light. Joshi also teaches that "photo oxidation of dioxyguanosine by riboflavin and UV radiation is of significant importance from the point of view of cell- damaging reactions by activated oxygen species produced by the synergistic action of sunlight and chemical agents. It is now known that activated oxygen species are responsible for skin photosensitization, tumor promotion and carcinogenic properties." Combining the teachings of Goodrich with the teachings of Joshi, one skilled in the art would think that irradiating blood products with riboflavin and UVB light would produce activated oxygen species which would cause damage to the red blood cells, platelets and plasma, and cause tumor promotion and cancer in the irradiated cells. Furthermore, as neither Goodrich nor Joshi teach the irradiation of white blood cells with UVB light, one skilled in the art would not think to do this using the combined teachings of Goodrich and Joshi.

Applicants' arguments have been fully considered but are not deemed persuasive for the following reasons. Goodrich et al. teach irradiating UV light to blood or blood components comprising red blood cells, white blood cells, platelets, plasma, bacteria and/or virus in the presence of riboflavin acting as a photosensitizer. Joshi teaches that riboflavin can be activated by exposing to UV-A (320-400 nm) and UV-B (290-320 nm) light. As such, one of ordinary skill in the art would have been motivated to use the methods taught by Goodrich et al. and Joshi in order to inactivate [1] donor

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white blood cells which can cause a series of severe immune responses in a transfusion recipient (as evidenced in Lee et al., From leukocyte reduction to leukocyte transfusion: the immunological effects of transfused leukocytes, Bailliere's Clinical Haematology, Vol. 13, No. 4, pp: 585-600, 2000), and [2] bacteria, viruses, and parasites which are potential sources of infection that have been transmitted by allogeneic transfusions. It is emphasized by the Examiner that in an obviousness-type rejection a single reference does not have to teach all the elements of the claims as long as the combined teachings of the prior art references meet the limitations of the claims. In this case, there is a clear motivation for combining the references of Goodrich et al. and Joshi as explained herein and in the previous office action.

Furthermore, it would have been obvious for one of skill in the art NOT to irradiate blood products, such as red blood cells and platelets, with an extreme concentration/dose of riboflavin and UVB light so that the irradiated cells become cancerous because a skilled artisan would have known the effect of extreme exposure to UVB light, and would have characterized the dose-response relationship of the amount of UVB irradiation cells are receiving and its effect (inactivation versus carcinogenesis). Likewise, it would have been obvious to a skilled artisan to characterize the dose-response relationship for the inactivation of white blood cells in order to determine the minimum dose of UVB required for activating riboflavin which results in the inactivation of white blood cells. As evidenced in the reference of Goodrich et al., the characterization of the dose-response relationship for irradiating cells with UV light in the presence of riboflavin was routine in the prior art with a finite

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number of predictable outcomes (see Examples 5 and 6). Therefore, for the reasons described herein and in the previous office action, the claimed invention as a whole is *prima facie* obvious over the combined teachings of the prior art.

Conclusion

Claims 12-15, 17-19 and 22 are rejected for the reasons as stated above.

Applicants must respond to the objections/rejections in this Office action to be fully responsive in prosecution.

THIS ACTION IS MADE FINAL. Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Jae W. Lee whose telephone number is 571-272-9949. The examiner can normally be reached on M-F between 9:00-6:00.

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If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Andrew Wang can be reached on 571-272-0811. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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/JAE W LEE/
Examiner, Art Unit 1656

/SUZANNE M. NOAKES/
Primary Examiner, Art Unit 1656